

estimated for an average respondent to respond: COPS Distressed Neighborhood Process Evaluation Survey: Eighteen respondents, at 1.5 hours per respondent (including record-keeping).

(6) An estimate of the total public burden (in hours) associated with the collection: Approximately 27 hours. If additional information is required contact: Ms. Brenda E. Dyer, Deputy Clearance Officer, United States Department of Justice, Information Management and Security Staff, Justice Management Division, Suite 850, Washington Center, 1001 G Street, NW, Washington, DC 20530.

Dated: June 23, 1999.

**Brenda E. Dyer,**

*Department Deputy Clearance Officer, United States Department of Justice.*

[FR Doc. 99-16445 Filed 6-28-99; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 21, 1999, Applied Science Labs, Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methcathinone (1237) .....	I
N-Ethylamphetamine (1475) .....	I
N,N-Dimethylamphetamine (1480).	I
4-Methylaminorex (cis isomer) (1590).	I
Lysergic acid diethylamide (7315)	I
Mescaline (7381) .....	I
3,4-Methylenedioxyamphetamine (7400).	I
N-Hydroxy-3,4-methylenedioxyamphetamine (7402).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxymethamphetamine (7405).	I
N-Ethyl-1-phenylcyclohexylamine (7455).	I
1-(1-Phenylcyclohexyl) pyrrolidine (7458).	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470).	I
Dihydromorphine (9145) .....	I

Drug	Schedule
Normorphine (9313) .....	I
Phenylcyclohexylamine (7460) .....	II
Phencyclidine (7471) .....	II
Phenylacetone (8501) .....	II
1-Piperidinocyclohexanecarbonitrile (8603).	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Benzoylcegonine (9180) .....	II
Morphine (9300) .....	II
Noroxymorphone (9668) .....	II

The firm plans to manufacture small quantities of the listed controlled substances for reference standards.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (60 days from publication).

Dated: June 22, 1999.

**John H. King,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 99-16415 Filed 6-28-99; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 27, 1999, Chiragene, Inc., 7 Powder Horn Drive, Warren, New Jersey 07059, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
N-Ethylamphetamine (1475) .....	I
2, 5-Dimethoxyamphetamine (7396).	I
3, 4-Methylenedioxyamphetamine (7400).	I
4-Methoxyamphetamine (7411) ...	I
Amphetamine (1100) .....	II
Methylphenidate (1724) .....	II

The firm plans to manufacture the listed controlled substances to supply their customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (60 days from publication).

Dated: June 22, 1999.

**John H. King,**

*Deputy Assistant Administrator Office of Diversion Control Drug Enforcement Administration.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importation of Controlled Substances; Notice of Application

Pursuant to section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on April 27, 1999, Radian International LLC, 14050 Summit Drive #121, P.O. Box 201088, Austin, Texas 78720-1088, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Cathinone (1235) .....	I
Methcathinone (1237) .....	I
N-Ethylamphetamine (1475) .....	I
l-bogaine (7260) .....	I
4-Bromo-2, 5-dimethoxyamphetamine (7391).	I
4-Bromo-2, 5-dimethoxyphenethylamine (7392).	I